## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## What is claimed is:

1. (Original) A method of preparing an optically pure compound having formula 1 or its salts, comprising:

reacting (R)-2-(4-methoxy-3-aminosulfonyl-phenyl)-1-methylethylamine or its salts with a compound selected from the group consisting of chloroacetic acid, bromoacetic acid, fluoroacetic acid, iodoacetic acid,  $\alpha$ -halogenoacetic acid anhydride, and  $\alpha$ -halogenoacetyl halide in the presence of a base or an acylating agent:

$$MeO$$
 $H_2NO_2S$ 
 $N$ 
 $H$ 
 $MeO$ 
 $MeO$ 

wherein

X is halogen.

- 2. (Original) The method of claim 1, wherein the base is trialkylamine or an inorganic base.
- 3. (Original) The method of claim 1, wherein the acylating agent is N,N'-dicyclohexylcarbodiimide, N,N'-diisopropylcarbodiimide, N-[3-(dimethylaminopropyl)-N'-ethylcarbodiimide, bis(2-oxo-3-oxazolidinyl)phosphinic chloride, (benzotriazole-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate, O-benzotriazole-1-yl-N,N,N',N'-bis(tetramethyl)uronium hexafluorophosphate, or O-(7-azabenzotriazole-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate.
- 4. (Original) The method of claim 1, wherein  $\alpha$ -halogenoacetic acid anhydride is obtained in-site during the reaction by reacting  $\alpha$ -halogenoacetic acid with at least one compound selected from the group consisting of methyl chloroformate, ethyl chloroformate, butyl chloroformate, benzyl chloroformate, and pivaloyl chloride.

- 5. (Original) The method of claim 1, wherein  $\alpha$ -halogenoacetyl halide is selected from the group consisting of bromoacetyl bromide, bromoacetyl chloride, chloroacetyl chloride, and chloroacetyl bromide.
- 6. (Currently Amended) A method of preparing (R)-5-{2-[2-(2-ethoxyphenoxy)ethylamino]-propyl}-2-methoxybenzenesulfonamide or its salts, comprising:

reacting an optically pure compound having formula 1 or its salts prepared using the method of any one-of claims 1 to 5 claim 1 with (i) 2-ethoxyphenol in the presence of a base or (ii) sodium 2-ethoxyphenoxide or potassium 2-ethoxyphenoxide to prepare (R)-2-(2-ethoxyphenoxy)-N-[2-(4-methoxy-3-aminosulfonyl-phenyl)- 1-methyl-ethyl]acetamide; and

reducing the obtained (R)-2-(2-ethoxyphenoxy)-N-[2-(4-methoxy-3-aminosulfonyl-phenyl)-1-methylethyl]acetamide.